4. (Amended) The method of claim 1 wherein the vessel has at least [one flexible] two walls a^2 with one wall being more deformable than the other wall. (Amended) The method of claim 1 wherein each of the particles [having] has a [mean] 6. volume of between about 5 x 10^{-24} m³ to about 5 x 10^{-6} m³. 7. (Amended) The method of claim 1 wherein the [substantial binding of the] particles [results at least in part from the particles having a coating] are coated. (Amended) The method of claim 1 wherein the coating comprises a polycationic 10. polymer. 12. (Amended) The method of claim 1 wherein the [network comprises] particles comprise a primary antibody and the additive comprises a secondary antibody, [where] the primary antibody [has] having a substantial binding to [the] a surface component of the cells, and the secondary antibody [has] having a substantial binding to the primary antibody. 13. (Amended) The method of [claim] any of claims 1 - 12 wherein the cells predominantly aś comprise red blood cells. 14. (Amended) The method of [claim] any of claims 1-12 wherein the [sample includes] blood cells comprise white blood cells and platelets. 15. (Amended) The method of any of claims 1 - 12, further comprising measuring [PSA] prostate specific antigen. 16. (Added) The method of claim 1, further comprising measuring creatinine. 17. (Amended) The method of any of claims 1 - 12 wherein at least 70% by volume of the [theoretically available] cell depleted portion is separated from the network within ten minutes. (Amended) The method of any of claims 1 - 12 wherein at least 90% by volume of the 21. [theoretically available] cell depleted portion is separated from the network within ten

minutes, with a separation efficiency of at least 95%.